Examiner: Marjorie A. Moran Group Art Unit: 1631

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A method for forming a two-dimensional ordered array of proteins, comprising:

contacting a population of proteins with a gas-aqueous interface;
laterally compressing said population to an appropriate pressure, such that
a two-dimensional ordered array of said proteins is formed at said interface, wherein said
proteins are not solubilized using detergent.

- 2. (Cancelled).
- 3. (**Previously Presented**) The method of claim 64, wherein said amphiphilic molecule comprises a protein.
- 4. (**Previously Presented**) The method of claim 1, wherein said protein is a membrane protein, a cellular receptor, an orphan receptor, receptor tyrosine kinase, an EPH receptor, an ion channel, a cytokine receptor, an multisubunit immune recognition receptor, a chemokine receptor, a growth factor receptor, or a G-protein coupled receptor.
- 5. (Currently Amended) The method of claim 1, wherein said protein is contacted with said interface in the presence of lipids.
- 6. (Currently Amended) The method of claim 1, further comprising applying said proteins to said interface in proteoliposomes, liposomes, or a cellular membrane.
- 7. (Cancelled).
- 8. (Currently Amended) The method of claim 1, wherein said interface is an airaqueous interface.

Claims 9-62 (Cancelled).

U.S.S.N. 10/003,468 Attorney Docket No. FMI-001RCE2

63. (Currently Amended) A method for forming a two- or three-dimensional ordered array of water insoluble membrane proteins, comprising:

Examiner: Marjorie A. Moran

Group Art Unit: 1631

contacting a population of <u>water insoluble</u> membrane proteins with a gasaqueous interface, wherein said population of membrane proteins are applied to said interface in a proteoliposome;

laterally compressing said population to an appropriate pressure, such that a two- or three-dimensional ordered array of said <u>water insoluble</u> membrane proteins is formed at said gas-aqueous interface.

64. (Currently Amended) A method for forming a three-dimensional ordered array of amphiphilic molecules water insoluble membrane proteins, comprising:

contacting a population of <u>amphiphilic molecules</u> <u>water insoluble</u> <u>membrane proteins</u> with a gas-aqueous interface;

laterally compressing said population to an appropriate pressure, such that a three-dimensional ordered array of said amphiphilic molecules water insoluble membrane proteins is formed at said interface, wherein said appropriate pressure is above a critical density point for the formation of a two-dimensional ordered array of said amphiphilic molecules water insoluble membrane proteins molecules.

Claims 65-66. (Cancelled).

- 67. (**Previously Presented**) The method of claim 1, wherein said two-dimensional ordered array is a two-dimensional crystalline array.
- 68. (**Previously Presented**) The method of claim 64, wherein said three-dimensional ordered array is a three-dimensional crystalline array.
- 69. (**Previously Presented**) The method of claim 3, wherein said protein is a membrane protein, a cellular receptor, an orphan receptor, receptor tyrosine kinase, an EPH receptor, an ion channel, a cytokine receptor, an multisubunit immune recognition receptor, a chemokine receptor, a growth factor receptor, or a G-protein coupled receptor.
- 70. (Previously Presented) The method of claim 3, wherein said protein is contacted with said interface in the presence of lipids.

U.S.S.N. 10/003,468 Attorney Docket No. FMI-001RCE2

71. (**Previously Presented**) The method of claim 3, further comprising applying said proteins to said interface in proteoliposomes, liposomes, or a cellular membrane.

Examiner: Marjorie A. Moran

Group Art Unit: 1631

Claims 72-73 (Cancelled).

74. (New) A method for forming a two- or three- dimensional ordered array of proteins suitable for use in crystallography to determine said protein's structure, comprising:

contacting a population of proteins with a gas-aqueous interface; laterally compressing said population to an appropriate pressure, such that a two-dimensional ordered array of said proteins is formed at said interface, wherein the structure of said protein using said two- or three- dimensional ordered array can be determined to a resolution of 5 Å or higher.

75. (New) A method for forming a two-dimensional ordered array of proteins, comprising:

contacting a population of proteins with a gas-aqueous interface;
laterally compressing said population to an appropriate pressure, such that
a two-dimensional ordered array of said proteins is formed at said interface, wherein said
two-dimensional ordered array is formed in the absence of a ligand of said protein.

76. (New) A method for forming a two- or three-dimensional ordered array of water insoluble membrane proteins, comprising:

contacting a population of water insoluble membrane proteins with a gasaqueous interface;

laterally compressing said population to an appropriate pressure, such that a two- or three-dimensional ordered array of said water insoluble membrane proteins is formed at said gas-aqueous interface.